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- (54) **METHOD OF SCREENING BAFF SUPPRESSOR OR INHIBITOR**
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- (52) **U.S. Cl.** **435/6; 435/7.23**
- (58) **Field of Classification Search** None
See application file for complete search history.

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(57) **ABSTRACT**

A method of screening a novel BAFF suppressor or inhibitor. More specifically speaking, a method which comprises adding a combination of TPA with ionomycin and/or an anti-CD3 antibody to a cultured human cell to thereby induce the production of BAFF by the cell; a method of screening a substance capable of suppressing the expression or activity of BAFF which comprises adding a test substance to a BAFF-production system prepared by adding a combination of TPA with ionomycin and/or an anti-CD3 antibody to a cultured human cell and measuring the expression amount and/or the activity of BAFF in the BAFF-production system; and a BAFF production inducer for a BAFF-producing cell which contains a combination of TPA with ionomycin and/or an anti-CD3 antibody.

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FIG. 1

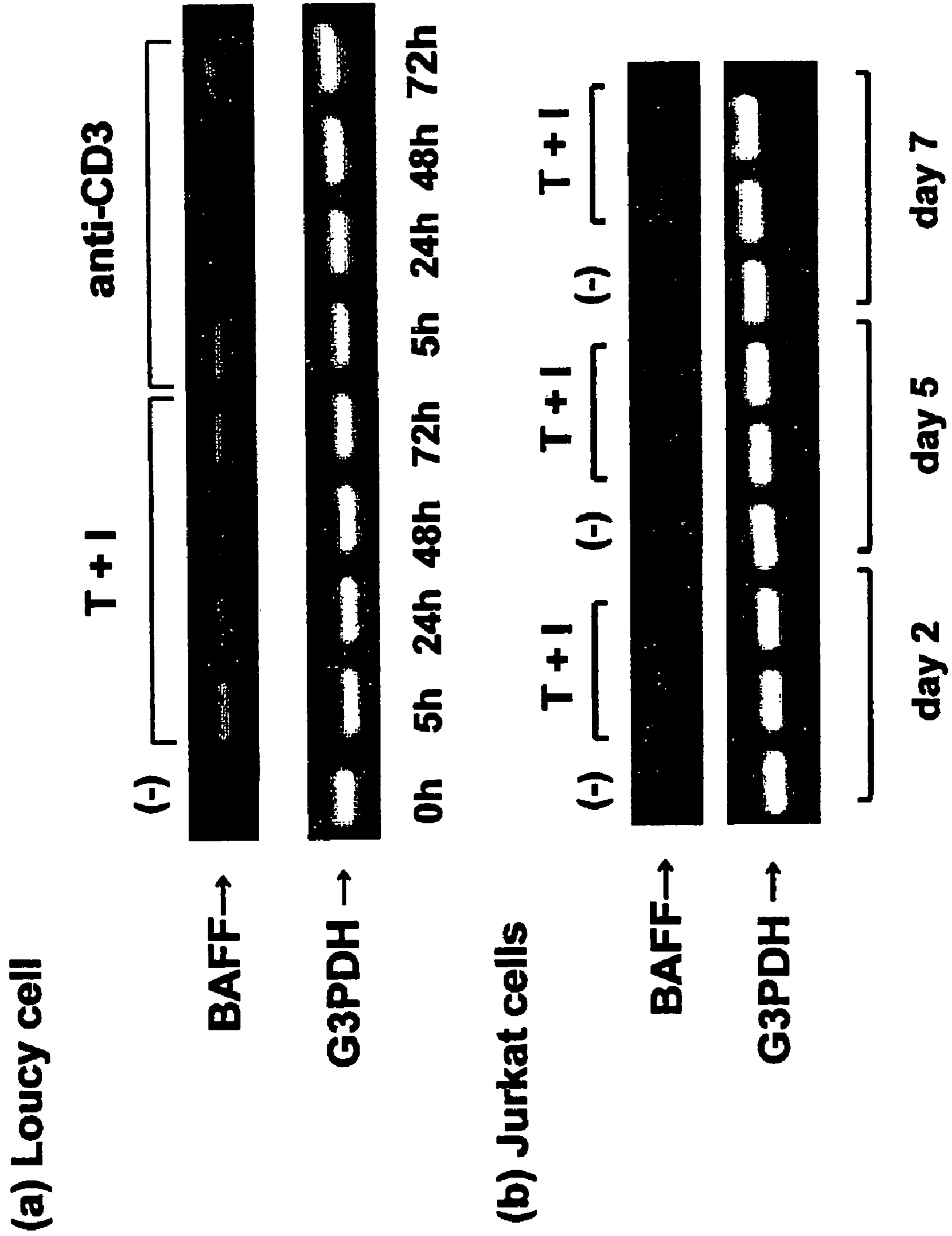


FIG. 2

Induction of BAFF in Loucy upon stimulation

CD3 stimulation TPA+Ionomycin

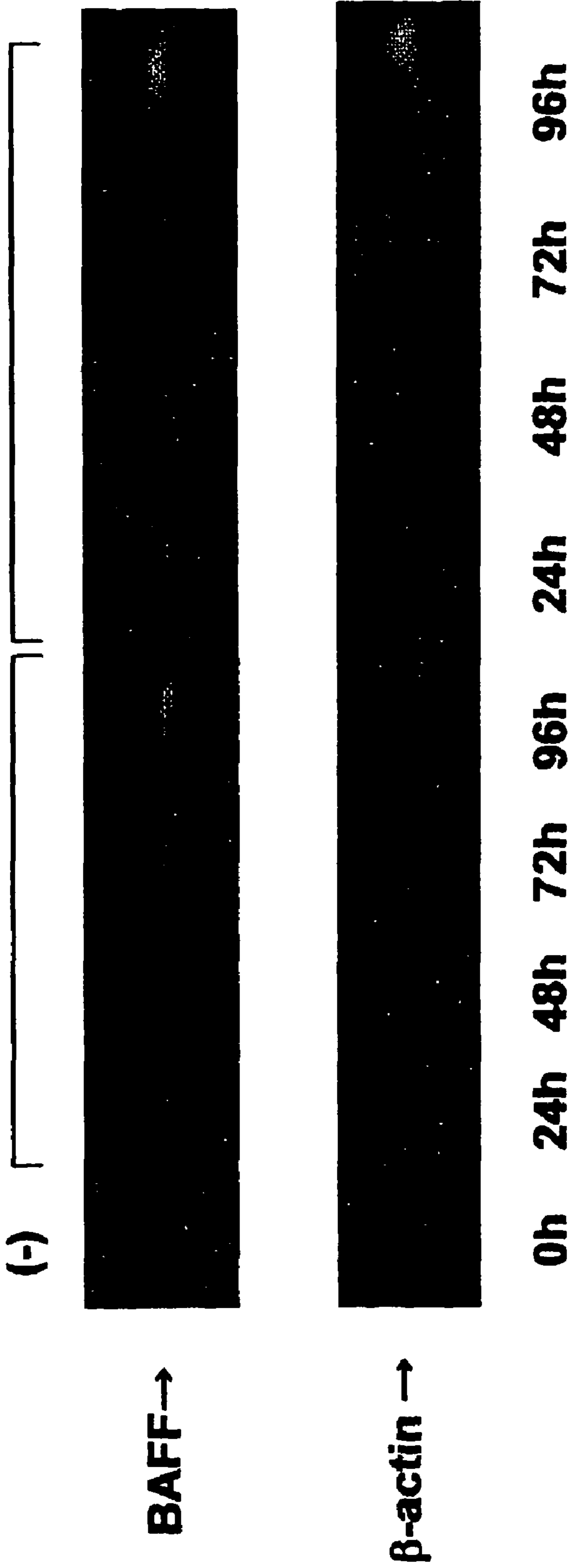
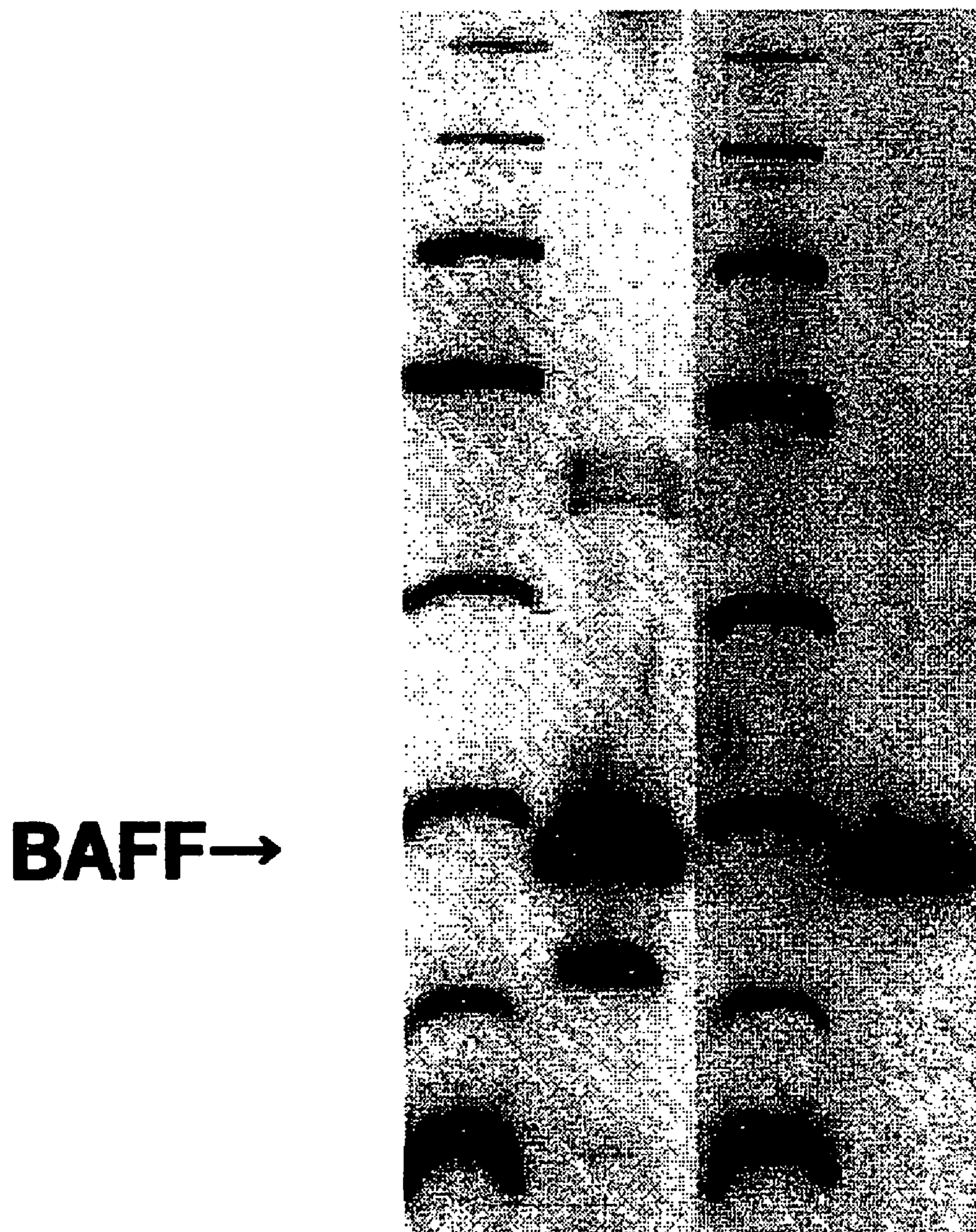


FIG. 3



Chem 4H4

FIG. 4

Production of BAFF from Loucy stimulated with TPA and Ionomycin

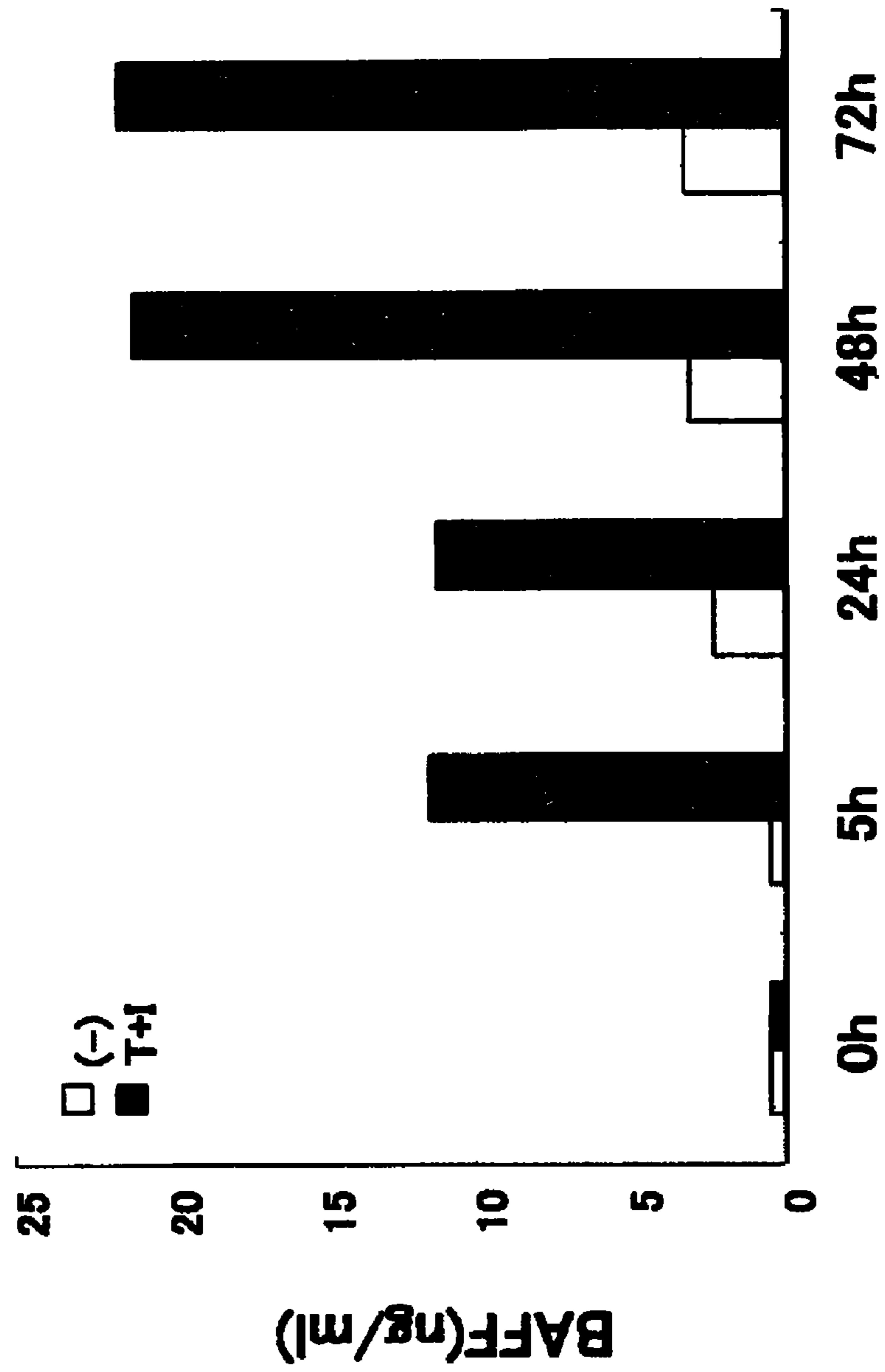
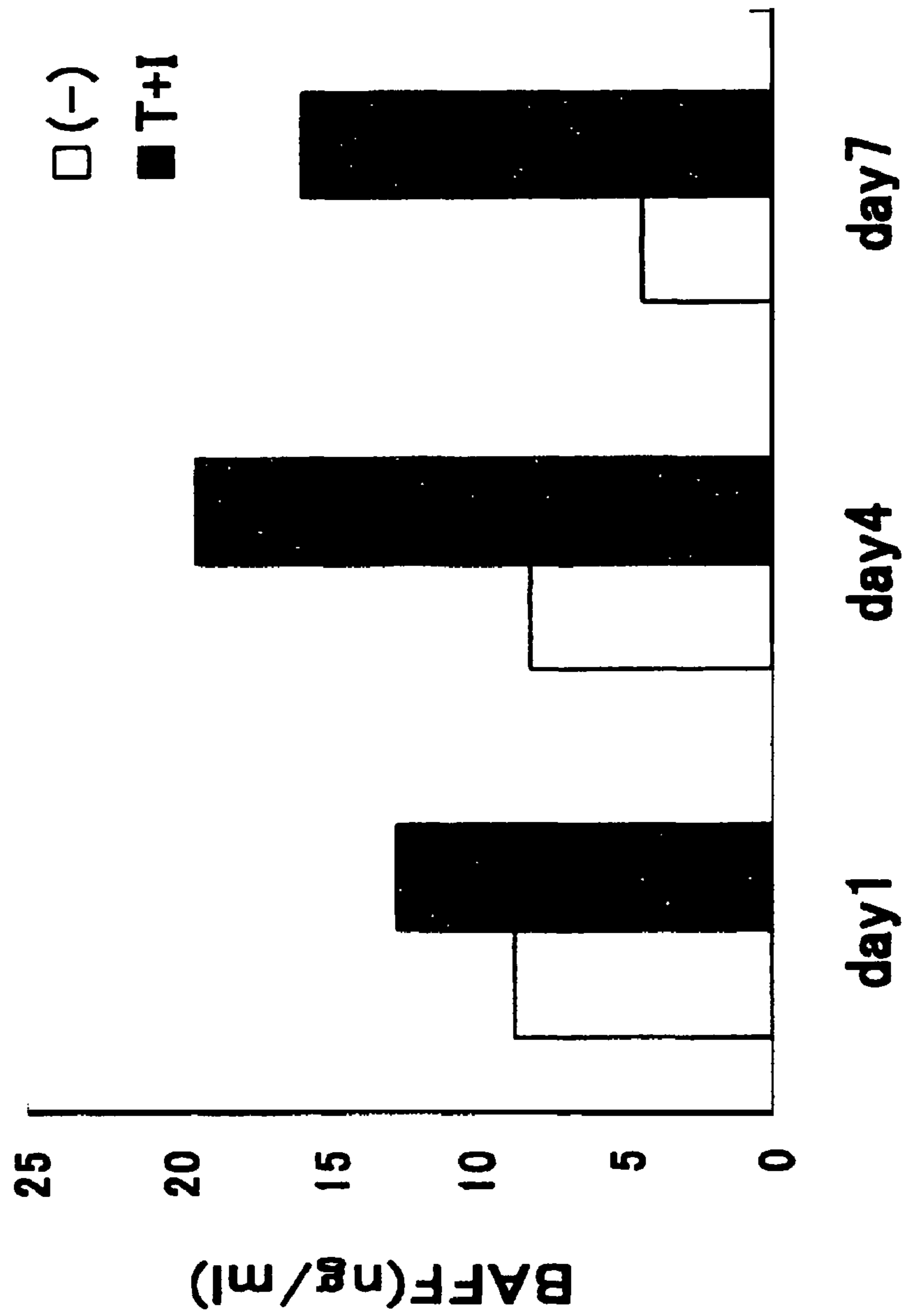


FIG. 5

Production of BAFF from Jurkat cells



1

METHOD OF SCREENING BAFF SUPPRESSOR OR INHIBITOR

TECHNICAL FIELD

The present invention relates to a method of screening a novel BAFF suppressor or inhibitor. The present invention relates particularly to a method of screening a substance suppressing or inhibiting the expression or production of BAFF by constructing a system wherein a detectable sufficient amount of BAFF is stably released extracellularly by subjecting a commercial cell strain to suitable stimulating conditions. Specifically, the present invention relates to a method of inducing the production of BAFF in a cultured human T cell by adding a combination of TPA with ionomycin and/or an anti-CD3 antibody to the cell, a method of screening a substance capable of regulating the expression or activity of BAFF, which includes adding a test substance to a BAFF production system in which a combination of TPA with ionomycin and/or an anti-CD3 antibody are/is added to a cultured human T cell and then measuring the expression level and/or activity level of BAFF in the BAFF production system, and a BAFF production inducer for a BAFF-producing cell, which comprises a combination of TPA with ionomycin and/or an anti-CD3 antibody.

BACKGROUND ART

It is known that BAFF (B cell activating factor belonging to the TNF family) is produced and secreted from T cells, monocytes/macrophages, dendritic cells and the like and regulates such as B-cell differentiation, activation, survival rate via 3 types of receptors on B cells (Moore et al., *Science*, 285, 260-263 (1999)).

Human BAFF is a transmembrane form protein comprising 285 amino acids. There is a structural characteristic of trimer formation such as the presence of a cytoplasmic domain of 46 amino acids, an extracellular domain of 218 amino acids, and two N-glycosylation sites in its amino acid sequence. It is estimated that an extracellular domain of 152 amino acids from C-terminal is cleaved with a protease of Furin family and released in a soluble form. The amino acid sequence of human BAFF initially named neutrokinine α was disclosed as SEQ ID NO: 1 or 2 in publication of International Patent Application WO98/18921. Other names of human BAFF such as Kay, TNFSF13B, Blys, TALL-1, THANK and zTNF4 are also known.

BAFF-R, TACI (transmembrane activator and calcium modulator and cyclophilin ligand interactor), and BCMA (B cell maturation antigen) are known as BAFF receptors. BAFF-R and BCMA are expressed mainly in B cells, and TACI is expressed in B cells and activated T cells.

The physiological action of BAFF lies in regulation of B-cell differentiation, activation, survival rate and the like as described above, and is increasingly revealed in recent years to participate in pathologic condition. That is, it is reported that a mouse expressing BAFF in excess shows SLE-like symptoms such as increase in peripheral blood B cells, enlargement of lymph nodes and spleen, increase in IgG level in serum, antinuclear antibody production, deposition of immune complex in the kidney, albuminuria and nephritis (Mackay et al., *J. Exp. Med.*, 190, 1697-1710, (1999), and Khare et al., *Proc. Natl. Acad. Sci. USA* 97, 3370-3375, (2000)). It was further revealed that this mouse also shows SS-like symptoms such as inflammation of salivary gland and destruction of salivary gland with advancing age (Groom et al., *J. Clin. Invest.*, 109, 59-68, (2002)) An increase of BAFF

2

level in serum in patients suffering from SLE, RA and SS is also reported (Zhanget al., *J. Immunol.*, 166, 6-10, (2001); Cheema et al., *Arthritis Rheum.*, 44, 1313-1319, (2001); and Groom et al., *J. Clin. Invest.*, 109, 59-68, (2002)), and there are also many reports such as higher BAFF level in synovial fluid than in serum in patients suffering from RA (Cheema et al., *Arthritis Rheum.*, 44, 1313-1319, (2001)), expression of BAFF in salivary gland-infiltrating leukocytes in patients suffering from SS (Groom et al., *J. Clin. Invest.* 109, 59-68, (2002)), correlation between serum BAFF level in patients suffering from SLE and immunoglobulin or anti-ds DNA antibody (Zhang et al., *J. Immunol.* 166, 6-10, (2001)) and correlation between BAFF in patients suffering from RA and rheumatoid factor (Cheema et al., *Arthritis Rheum.*, 44, 1313-1319, (2001)).

From these facts, it can be said that the suppression or inhibition of expression or production of BAFF leads to prevention and treatment of autoimmune diseases such as SLE, RA and SS, and thus there is demand for establishment of a system for accurately screening the suppression or inhibition of expression or production of BAFF.

For establishment of such screening system, it is possible to anticipate, for example, a method of obtaining a cell expressing BAFF stably by transforming a cultured cell with a gene vector constructed so as to express BAFF by recombinant DNA techniques. There is however a problem that even if BAFF could be expressed, BAFF would not always be released extracellularly in a secretory form and would be uncertain as to whether the cultured cell can be a cell strain stable for expression and production of BAFF. Accordingly, none of such system has been reported. For example, if it would be possible to construct a system wherein the induction of intracellular expression of BAFF and the extracellular release of a detectable sufficient amount of BAFF can be recognized when a commercial cell strain is subjected to suitable stimulating conditions, the problem described above could be solved.

DISCLOSURE OF INVENTION

The object of the present invention is to provide a method of screening a substance suppressing or inhibiting the expression or production of BAFF by constructing a system wherein a detectable sufficient amount of BAFF is stably released extracellularly by subjecting a commercial cell strain to suitable stimulating conditions. By constructing such system, there can also be provided a method of screening a prophylactic/therapeutic agent for autoimmune diseases such as systemic lupus erythematosus (SLE), chronic rheumatoid arthritis (RA), Sjogren's syndrome (SS), autoimmune diabetes, AIDS, and an autoimmune disease accompanied by B-cell activation.

Accordingly, the present inventors made extensive study on selection of a cell strain and a method of stimulating the cell, and as a result, found that when a Loucy cell or Jurkat cell that is a commercial human T cell strain highly expressing CD3 is stimulated with an anti-CD3 antibody and/or a combination of TPA (12-O-tetradecanoylphorbol 13-acetate, or Phorbol 12-myristate 13-acetate, also expressed as PMA) with ionomycin, the induction of stable BAFF expression in the cell is recognized and simultaneously a detectable sufficient amount of BAFF can be stably released into a culture, and the present invention was thereby completed.

That is, the present invention provides a BAFF production system wherein a cultured human T cell is stimulated by adding a combination of TPA with ionomycin and/or an anti-CD3 antibody to the cell. Specifically, the present invention

3

provides a BAFF production system wherein a cultured human T cell such as Loucy cell or Jurkat cell is stimulated by adding a combination of TPA with ionomycin and/or an anti-CD3 antibody to the cell.

The present invention also relates to a method of inducing the production of BAFF in a cultured human T cell, by adding a combination of TPA with ionomycin and/or an anti-CD3 antibody to the cell. Specifically, the present invention relates to a method of inducing the production of BAFF in a cultured human T cell such as Loucy cell or Jurkat cell, by adding a combination of TPA with ionomycin and/or an anti-CD3 antibody to the cell.

The present invention also provides a method of screening a substance capable of regulating the expression or activity of BAFF in a BAFF production system wherein a cultured human T cell is stimulated by adding a combination of TPA with ionomycin and/or an anti-CD3 antibody to the cell. Specifically, the present invention provides a method of screening a substance capable of suppressing or inhibiting the expression or activity of BAFF in a BAFF production system wherein a cultured human T cell is stimulated by adding a combination of TPA with ionomycin and/or an anti-CD3 antibody to the cell. More specifically, the present invention provides a method of screening a substance capable of regulating, preferably suppressing or inhibiting the expression or activity of BAFF in a BAFF production system wherein a cultured human T cell such as Loucy cell or Jurkat cell is stimulated by adding a combination of TPA with ionomycin and/or an anti-CD3 antibody to the cell.

The screening method of the present invention relates particularly to a method of screening a substance capable of regulating the expression or activity of BAFF, which includes adding a test substance to a BAFF production system in which a combination of TPA with ionomycin and/or an anti-CD3 antibody are/is added to a cultured human T cell and then measuring the expression level and/or activity level of BAFF in the BAFF production system. Specifically, the present invention relates to a method of screening a substance capable of regulating the expression or activity of BAFF, which includes adding a test substance to a BAFF production system in which a combination of TPA with ionomycin and/or an anti-CD3 antibody are/is added to a cultured human T cell such as Loucy cell or Jurkat cell and then measuring the expression level and/or activity level of BAFF in the BAFF production system. More specifically, the present invention relates to a method of screening a substance capable of suppressing or inhibiting the expression or activity of BAFF, which includes adding a test substance to a BAFF production system in which a combination of TPA with ionomycin and/or an anti-CD3 antibody are/is added to a cultured human T cell such as Loucy cell or Jurkat cell and then measuring the expression level and/or activity level of BAFF in the BAFF production system.

Further, the present invention provides a method of screening a prophylactic/therapeutic agent for autoimmune diseases such as systemic lupus erythematosus (SLE), chronic rheumatoid arthritis (RA), Sjogren's syndrome (SS), autoimmune diabetes, AIDS, and an autoimmune disease accompanied by B-cell activation, in a BAFF production system wherein a cultured human T cell is stimulated by adding a combination of TPA with ionomycin and/or an anti-CD3 antibody to the cell. Specifically, the present invention provides a method of screening a prophylactic/therapeutic agent for autoimmune diseases such as systemic lupus erythematosus (SLE), chronic rheumatoid arthritis (RA), Sjogren's syndrome (SS), autoimmune diabetes, AIDS, and an autoimmune disease accompanied by B-cell activation, in a BAFF production

4

system wherein a cultured human T cell such as Loucy cell or Jurkat cell is stimulated by adding an anti-CD3 antibody and/or a combination of TPA with ionomycin to the cell. More specifically, the present invention provides a method of screening a prophylactic/therapeutic agent for autoimmune diseases such as systemic lupus erythematosus (SLE), chronic rheumatoid arthritis (RA), Sjogren's syndrome (SS), autoimmune diabetes, AIDS, and an autoimmune disease accompanied by B-cell activation, which includes adding a test substance to a BAFF production system in which a combination of TPA with ionomycin and/or an anti-CD3 antibody are/is added to a cultured human T cell such as Loucy cell or Jurkat cell and then measuring the expression level and/or activity of BAFF in the BAFF production system.

Further, the present invention relates to a BAFF production inducer for a BAFF-producing cell, which contains a combination of TPA with ionomycin and/or an anti-CD3 antibody. Specifically, the present invention relates to a BAFF production inducer for a cultured human T cell, which contains a combination of TPA with ionomycin and/or an anti-CD3 antibody. More specifically, the present invention relates to a BAFF production inducer for a cultured human T cell such as Loucy cell or Jurkat cell, which contains a combination of TPA with ionomycin and/or an anti-CD3 antibody.

According to the present invention, there can be provided a method of screening, with a cultured cell strain, a substance capable of suppressing or inhibiting the production of BAFF as well as a method of screening a prophylactic/therapeutic agent for diseases involved in production or activity of BAFF, such as systemic lupus erythematosus (SLE), chronic rheumatoid arthritis (RA), Sjogren's syndrome (SS), autoimmune diabetes, AIDS, and an autoimmune disease accompanied by B-cell activation.

BRIEF DESCRIPTION OF DRAWINGS

FIG. 1 is a drawing showing the expression of BAFF upon stimulation of a Loucy cell or Jurkat cell with an anti-CD3 antibody or both TPA and ionomycin.

FIG. 2 is a drawing showing the expression of BAFF protein upon stimulation of a Loucy cell with an anti-CD3 antibody or both TPA and ionomycin.

FIG. 3 is a drawing in which recombinant human BAFF (manufactured by Chemicon) was detected by conventional Western blotting method with an antihuman BAFF monoclonal antibody (4H4) and a control rabbit antihuman BAFF polyclonal antibody (Chem) manufactured by Chemicon.

FIG. 4 is a graph showing the production of BAFF upon stimulation of a Loucy cell with TPA and ionomycin.

FIG. 5 is a graph showing the production of BAFF upon stimulation of a Jurkat cell with TPA and ionomycin.

BEST MODE FOR CARRYING OUT THE INVENTION

The inventors examined the expression of BAFF upon stimulation of a Loucy cell or Jurkat cell with various substances. As a result, the inventors found that when a Loucy cell or Jurkat cell is stimulated with an anti-CD3 antibody or a combination of TPA with ionomycin, the induction of stable BAFF expression in the cell is recognized and simultaneously a detectable sufficient amount of BAFF can be stably released into a culture, as shown later in the Examples.

Accordingly, the method of screening a substance capable of regulating, preferably suppressing or inhibiting the expression or activity of BAFF can be constituted as follows.

That is, commercial human T cells are suspended at a suitable density in a suitable medium, then seeded in a culture plate or the like, and stimulated by adding TPA (final concentration of 0.001 to 1.0 $\mu\text{g}/\text{mL}$, preferably 0.005 to 0.1 $\mu\text{g}/\text{mL}$) and ionomycin (final concentration of 0.01 to 10.0 $\mu\text{g}/\text{mL}$, preferably 0.05 to 1.0 $\mu\text{g}/\text{mL}$), and a test substance (any concentration gradients at a final concentration of 10^{-15} to 10^{-3} M) is added as necessary, and culture of the cells is initiated. The culture supernatant is recovered 1 hour to 7 days, preferably 3 hours to 5 days, after the culture is initiated, and the concentration of BAFF in the supernatant can be measured and compared with the control, thereby measuring the activity of the test substance to suppress or inhibit BAFF in this screening system. For carrying out the above-described system, a culture plate coated previously with an anti-CD3 antibody diluted with PBS or the like (final concentration of 0.01 to 1000 $\mu\text{g}/\text{mL}$, preferably 0.1 to 100 $\mu\text{g}/\text{mL}$) may be used. The culture plate or the like coated with an anti-CD3 antibody can be used to detect a sufficient amount of BAFF without using TPA and ionomycin and is useful as an easier screening system.

The human T cell strain used in the present invention is preferably a commercial cell, and examples include a Loucy cell (ATCC Cat. No. CRL-2629), a Jurkat cell (ATCC Cat. No. TIB-152) and a TALL-1 cell (JCRB 0086), among which the Loucy cell (ATCC Cat. No. CRL-2629) or Jurkat cell (ATCC Cat. No. TIB-152) highly expressing CD3 and expected to be activated via a T-cell receptor is more preferable.

The anti-CD3 antibody used in the present invention includes an antihuman CD3 monoclonal antibody and antihuman CD3 polyclonal antibody produced by methods known by those skilled in the art, among which an antihuman CD3 monoclonal antibody is preferable, and a mouse antihuman CD3 monoclonal antibody (BD Pharmingen Cat No. 555336, or the like) is more preferable.

TPA used in the present invention, which is an abbreviation of 12-O-tetradecanoylphorbol 13-acetate, is a phorbol ester also known as phorbol 12-myristate 13-acetate (abbreviated sometimes as PMA), and TPA used in the present invention may be any of compounds given such names and also includes stereoisomers. Preferable commercial TPA includes, for example, 4- α -phorbol 12-myristate 13-acetate (SIGMA Cat. No. P-148).

Ionomycin used in the present invention is a polyether antibiotic derived from *Streptomyces conglobatus* and is known to act as calcium ionophore, and ionomycin used in the present invention also includes pharmaceutically acceptable salts. Preferable commercially available ionomycin includes, for example, ionomycin, calcium salt (SIGMA Cat. No. IO634).

There is a characteristic that BAFF production inducer of the present invention contains a combination of TPA with ionomycin and/or an anti-CD3 antibody as active ingredients for inducing the production of BAFF. The BAFF production inducer of the present invention can comprise, in addition to the above active ingredients, a carrier such as a buffer solution which can be added to a cell system, and together with the carrier which can be added to a cell system, can be formed into a composition for inducing the production of BAFF.

The BAFF production inducer of the present invention can be applied to a BAFF-producing cell, preferably a cultured cell, and preferable examples of the cell include a cultured human T cell, specifically a Loucy cell and Jurkat cell.

When the BAFF production inducer of the present invention includes a combination of TPA with ionomycin as the active ingredients, the respective active ingredients are used

by adding them to a cell system in such amounts that the final concentration of TPA becomes 0.001 to 1.0 $\mu\text{g}/\text{mL}$, preferably 0.005 to 0.1 $\mu\text{g}/\text{mL}$, while the final concentration of ionomycin becomes 0.01 $\mu\text{g}/\text{mL}$ to 10 $\mu\text{g}/\text{mL}$, preferably 0.05 $\mu\text{g}/\text{mL}$ to 1 $\mu\text{g}/\text{mL}$. When the active ingredient is an anti-CD3 antibody, the active ingredient can be put to a culture plate at a concentration of 0.1 $\mu\text{g}/\text{mL}$ to 100 $\mu\text{g}/\text{mL}$, preferably 1 $\mu\text{g}/\text{mL}$ to 30 $\mu\text{g}/\text{mL}$ in a buffer solution.

The thus obtained BAFF production system of the present invention has a surprising feature that a detectable sufficient amount of BAFF can be stably released into a culture as shown later in the Examples, and thus it is possible to realize not only a method of screening a substance capable of regulating, preferably suppressing or inhibiting the expression or activity of BAFF, but also a method of screening a prophylactic/therapeutic agent for autoimmune diseases such as systemic lupus erythematosus (SLE), chronic rheumatoid arthritis (RA), Sjogren's syndrome (SS), autoimmune diabetes, AIDS, and an autoimmune disease accompanied by B-cell activation.

EXAMPLES

Hereinafter, the present invention is described in more detail by reference to the Examples, but the technical scope of the present invention is not limited by the Examples.

Example 1

Expression of BAFF upon stimulation of a Loucy cell or Jurkat Cell with an Anti-CD3 Antibody or Both TPA and Ionomycin

(a) Culture Under Stimulation with TPA and Ionomycin

Loucy cells or Jurkat cells purchased from ATCC (American Type Culture Collection) were suspended in RPMI1640 medium containing 10% FBS (Fetal bovine serum), 0.45% D-glucose, 1 mM sodium pyruvate and 1 mM HEPES and seeded at a density of 1×10^6 cells/well on a 6-well culture plate. At the same time, TPA and ionomycin were added respectively to a final concentration of 0.05 $\mu\text{g}/\text{mL}$, 0.5 $\mu\text{g}/\text{mL}$ or 0.1 $\mu\text{g}/\text{mL}$, and the cells were recovered 5 hours, 24 hours, 48 hours and 72 hours (Loucy cells) or 48 hours, 120 hours and 168 hours (Jurkat cells) after the culture was initiated. Total RNA was extracted from these recovered cells, and BAFF mRNA was analyzed by the RT-PCR method under the following conditions. Using ReverTra Dash kit (TOYOBO Cat. No. PCR-401), the sample was subjected to reverse transcription reaction at 30° C. for 10 minutes, at 42° C. for 20 minutes and at 99° C. for 5 minutes to give cDNA. Using Advantage 2 PCR Enzyme system (BD Biosciences Cat. No. PT3281-1), this cDNA was reacted at 94° C. for 1 minute, then subjected to 35 reaction cycles each at 94° C. for 30 seconds and at 66° C. for 1.5 minutes and reacted at 66° C. for 5 minutes, and the DNA thus acquired was separated by 1.5% agarose gel electrophoresis, and an objective band was recognized at a position of 975 bp. The primer sequences used in the PCR reaction are as follows:

Forward:
5'-aggccccaaccttcaaagttcaag (SEQ ID NO: 1)

Reverse:
5'-cttagaggtacagagaaagggagg (SEQ ID NO: 2)

The results are shown in FIG. 1. FIG. 1(a) shows the sample from the Loucy cells, and FIG. 1(b) shows the sample

from the Jurkat cells. T+I shows stimulation with TPA and ionomycin (which are 0.05 $\mu\text{g}/\text{mL}$ and 0.5 $\mu\text{g}/\text{mL}$ respectively in (a); and 0.05 $\mu\text{g}/\text{mL}$ and 0.5 $\mu\text{g}/\text{mL}$ (on the observers' right) or 0.1 $\mu\text{g}/\text{mL}$ (on the observers' left) in (b)), and (-) shows non-stimulation. In each profile, G3PDH is a positive control. In Loucy cells and Jurkat cells stimulated with TPA and ionomycin, the expression of BAFF mRNA was recognized 5 hours and 2 days after the stimulation, and the induction of evident expression of BAFF mRNA by TPA and ionomycin was continuously recognized over 72 hours and 7 days (168 hours) after the culture was initiated, as shown in FIGS. 1(a) and (b). It was thus found that because the induction of stable expression of BAFF was recognized by culturing Loucy cells or Jurkat cells under stimulation with TPA and ionomycin, this culture system can be used in screening a substance suppressing or inhibiting the expression of BAFF.

(b) Culture Under Stimulation with an Anti-CD3 Antibody

An anti-CD3 antibody diluted at 10 $\mu\text{g}/\text{mL}$ with PBS was put to a 6-well plate and adsorbed onto the bottom at 4° C. overnight, and then Loucy cells suspended in the same manner as in (a) were seeded at a density of 1×10^6 cells/well. The cells were recovered 5 hours, 24 hours, 48 hours and 72 hours after the culture was initiated. Total RNA was extracted from these recovered cells, and BAFF mRNA was analyzed by the RT-PCR method under the following conditions. Using ReverTra Dash kit (TOYOBO Cat. No. PCR-401), the sample was subjected to reverse transcription reaction at 30° C. for 10 minutes, at 42° C. for 20 minutes and at 99° C. for 5 minutes to give cDNA. Using Advantage 2 PCR Enzyme system (BD Biosciences Cat. No. PT3281-1), this cDNA was reacted at 94° C. for 1 minute, then subjected to 35 reaction cycles each at 94° C. for 30 seconds and at 66° C. for 1.5 minutes and reacted at 66° C. for 5 minutes, and the acquired DNA was separated by 1.5% agarose gel electrophoresis, and an objective band was recognized at a position of 975 bp. As a result, two-phase expression was made evident as shown in anti-CD3 in the right in FIG. 1(a), wherein the expression of BAFF mRNA is recognized from 5 hours after stimulation, thereafter the expression is attenuated, and the mRNA is expressed again 72 hours after the culture is initiated. There would be a possibility that such expression modes are different in biological meaning, and it is considered that the expression of BAFF induced by culturing Loucy cells under stimulation with an anti-CD3 antibody can be used in screening a substance suppressing or inhibiting the unique expression of BAFF.

Example 2

Expression of BAFF Protein Upon Stimulation of a Loucy Cell with an Anti-CD3 Antibody or Both TPA and Ionomycin

An anti-CD3 antibody diluted at 10 $\mu\text{g}/\text{mL}$ with PBS was put to a 6-well plate and adsorbed onto the bottom at 4° C. overnight, and then Loucy cells suspended in the same manner as in Example 1 were seeded at a density of 1×10^6 cells/well. The cells were recovered 24 hours, 48 hours, 72 hours and 96 hours after the culture was initiated. These recovered cells were used and examined for expression of BAFF protein by a conventional Western blotting method. As an antibody for detection of BAFF protein, a rabbit antihuman BAFF polyclonal antibody (AB16530, manufactured by Chemicon) was used. The results are shown in FIG. 2. A control not stimulated is shown in (-) in the left in FIG. 2, and CD3 stimulation in FIG. 2 shows stimulation with anti-CD3 anti-

body, and TPA+Ionomycin shows stimulation with TPA and ionomycin. β -actin in the lower profile in FIG. 2 shows a positive control. In the Loucy cells stimulated with an anti-CD3 antibody, a 34-kDa band corresponding to human BAFF was confirmed from 24 hours after stimulation, and this evident expression induction was continuously recognized over 96 hours after the culture was initiated. Separately, Loucy cells were suspended in RPMI1640 medium containing 10% FBS (fetal bovine serum), 0.45% D-glucose, 1 mM sodium pyruvate and 1 mM HEPES and seeded at a density of 1×10^6 cells/well on a 6-well culture plate, and at the same time, simultaneously TPA and ionomycin were added to final concentrations of 0.05 $\mu\text{g}/\text{mL}$ and 0.5 $\mu\text{g}/\text{mL}$ respectively, and the cells were recovered 24 hours, 48 hours, 72 hours and 96 hours after the culture was initiated. These recovered cells were used and examined for expression of BAFF protein by a conventional Western blotting method. In the Loucy cells stimulated with TPA and ionomycin, a 34-kDa band corresponding to human BAFF was recognized from 24 hours after stimulation, and this evident expression induction was continuously recognized over 96 hours after the culture was initiated, as shown in FIG. 2. In these results, the induction of stable expression of BAFF protein by culturing Loucy cells under stimulation with an anti-CD3 antibody or both TPA and ionomycin was recognized over 96 hours after the culture was initiated, and this culture system can be used in screening a substance suppressing or inhibiting the expression of BAFF protein.

Reference Example 1

Preparation of Antihuman BAFF Monoclonal Antibody (4H4)

The antihuman BAFF monoclonal antibody (4H4) used in Example 3 was prepared in the following manner. That is, 13 amino acids (SEQ ID NO: 4) corresponding to a region, in the vicinity of a membrane, of an extracellular domain in 285 amino acids of BAFF shown in SEQ NO: 3 in the Sequence Listing were selected, then conjugated with KLH by the MBS method, and used as antigen. 100 μL of 1 mg/ml aqueous solution of the antigen peptide in physiological saline and Freund's complete adjuvant were formed into an emulsion and then used in intraperitoneally immunizing a mouse (Balb/c, 6-week-old). After 2 weeks, 100 μL of 1 mg/ml of an aqueous solution of the antigen peptide in physiological saline and Freund's complete adjuvant, which had been emulsified by sonication, were used as booster for additional immunization, followed by additional immunization twice at 2-week intervals. Two months after the first immunization, the spleen was excised, and lymphocytes were separated in RPMI 1640 medium (containing penicillin and streptomycin). The separated lymphocytes were fused with mouse bone marrow-derived myeloma cell P3U1 strain by the polyethylene glycol (PEG) method to prepare hybridoma cells. The hybridoma cells were suspended in a feeder cell-containing HAT medium, pipetted to a 96-well plate (Greiner) and cultured for 15 days. A culture supernatant was recovered from the wells in which the hybridoma cells had been cultured, and antibody-producing cells reactive with the antigen peptide were selected by ELISA (enzyme-linked immunosorbent assay). That is, first, 50 μL of 10 $\mu\text{g}/\text{mL}$ antigen peptide was put to each well of the 96-well plate, adsorbed onto the bottom at 4° C. overnight and blocked with 100 μL of 2% BSA/PBS at 37° C. for 2 hours. Each well was reacted at 4° C. overnight with 100 μL supernatant of the hybridoma cells and then reacted at 37° C. for 1 hour with a 1000-fold dilution of

HRP-labeled anti-mouse IgG and colored with orthophenylene diamine as substrate. After the reaction was terminated with 50 μ L of 2 N sulfuric acid, each well was measured for absorption at 492 nm, and hybridomas showing an absorption of 1.0 or more were selected and cloned by limiting dilution.

The selected hybridoma cells were injected intraperitoneally to a mouse (Balb/c) to which 0.5 mL pristane had been intraperitoneally administered before 7 days and before 3 days, and about 10 days later, the ascites fluid was collected. The recovered ascites fluid was left at room temperature for 30 minutes, then left at 4° C. overnight, centrifuged at 15 Krpm for 10 minutes to recover a supernatant from which a mouse IgG fraction was separated and purified through a protein A-Sepharose column.

By the method described above, a hybridoma cell strain producing the antihuman BAFF antibody (4H4 whose isotype is IgG1), as well as the antibody (4H4), was obtained. The resulting 4H4, and a control rabbit antihuman BAFF polyclonal antibody (AB16530: shown as Chem in FIG. 3) manufactured by Chemicon, were used in detection of recombinant human BAFF (manufactured by chemicon) by the conventional Western blotting method. As a result, a 17-KDa band corresponding to the soluble human BAFF was confirmed as shown in FIG. 3.

Example 3

Production of BAFF Upon Stimulation of a Loucy Cell or Jurkat Cell with TPA and Ionomycin

Loucy cells or Jurkat cells suspended in the same manner as in Example 1 were seeded to a 6-well culture plate at a density of 1×10^6 cells/well. At the same time, TPA and ionomycin were added to final concentrations of 0.05 μ g/mL and 0.5 μ g/mL respectively, and the culture supernatant was recovered 5 hours, 24 hours, 48 hours, 72 hours (Loucy cells) or 1, 4 and 7 days (Jurkat cells) after the culture was initiated. BAFF produced by the cells contained in these culture supernatants was measured by an ELISA method using a commercial rabbit antihuman BAFF polyclonal antibody (Cat No. AB16530, manufactured by Chemicon) and the mouse antihuman BAFF monoclonal antibody (4H4) prepared in Reference Example 1. The ELISA method was carried out in the following manner. That is, a 96-well plate was coated at 4° C. overnight with 1 μ g/well rabbit antihuman BAFF polyclonal antibody (Cat. No. AB16530, manufactured by Chemicon) as primary antibody. Each well was washed 3 times with PBS

containing 0.05% Tween 20, and then Block Ace (Dainippon Pharmaceutical Co., Ltd.) was added in a volume of 150 μ L/well and reacted at 37° C. for 2 hours. Each well was washed 3 times with PBS containing 0.05% Tween 20, and 50 μ L sample and 50 μ L (8 ng/mL) of biotin-labeled 4H4 were added and reacted at room temperature for 2 hours. Each well was washed 3 times with PBS containing 0.05% Tween 20, and 50 μ L of a 1000-fold dilution of streptavidin-labeled HRP (Horse Radish peroxidase) diluted with PBS containing 0.05% Tween 20 was added and reacted at room temperature for 30 minutes. Each well was washed 5 times with PBS containing 0.05% Tween 20, and then 50 μ L of TMB One Solution (manufactured by Clonetech) was added and reacted for 5 minutes at room temperature, then 50 μ L of 1 N HCl was added, and the each well was measured for absorbance at 450 nm with a plate reader (manufactured by Perkin Elmer). As a standard substance, recombinant human BAFF (Cat No. GF119, manufactured by Chemicon) was used to prepare a standard curve, from which the concentration of BAFF in the sample was measured.

A graph showing the result where the Loucy cells were used is shown in FIG. 4, and a graph showing the result where the Jurkat cells were used is shown in FIG. 5. The concentration of BAFF (ng/mL) is shown on the ordinate in each graph, and each left bar graph shows, as control, the concentration of BAFF without stimulation with TPA and ionomycin. The results indicated, as shown in FIGS. 4 and 5, that in the Loucy cells or Jurkat cells stimulated with TPA and ionomycin, BAFF in the culture supernatant can be detected from 5 hours or 1 day after stimulation and is continuously detected over 72 hours or 7 days after the culture was initiated. From these results, it was revealed that the induction of stable expression of BAFF was recognized by culturing Loucy cells or Jurkat cells under stimulation with TPA and ionomycin, and this culture system can be used in screening a substance suppressing or inhibiting the production of BAFF.

INDUSTRIAL APPLICABILITY

The present invention provides a method of inducing, in a cultured human T cell, the production of BAFF revealed to be related to autoimmune diseases such as systemic lupus erythematosus (SLE), chronic rheumatoid arthritis (RA), Sjogren's syndrome (SS), autoimmune diabetes, AIDS, and an autoimmune disease accompanied by B-cell activation, and provides a technique of screening a new substance useful for treatment and prevention of autoimmune diseases, and is industrially applicable.

SEQUENCE LISTING

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Ser Phe Tyr Gln Val Ala Ala Leu Gln Gly Asp Leu Ala Ser Leu Arg
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Ser Leu Val Thr Leu Phe Arg Cys Ile Gln Asn Met Pro Glu Thr Leu
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Pro Asn Asn Ser Cys Tyr Ser Ala Gly Ile Ala Lys Leu Glu Glu Gly
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The invention claimed is:

1. A method of screening a substance capable of regulating the expression or activity of B cell activating factor belonging to the tumor necrosis factor family (BAFF), comprising:

adding a test substance to a BAFF production system including a Loucy cell stimulated by a combination of 12-O-tetradecanoylphorbol 13-acetate (TPA) and ionomycin, and then

measuring at least one of the expression level and activity level of BAFF in the BAFF production system.

2. The method according to claim 1, wherein the regulation¹⁵ of expression or activity of BAFF is the suppression or inhibition of expression or activity of BAFF.

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